Methods

The winegrape cuttings used in this experiment were taken from the UC Davis Robert Mondavi Institute vineyard in December (USE YEARS? European, introduce with year, then add if it changes). They were potted in 26 cm diameter pots and began growing in January. On May 27, they were placed in growth chambers with day/night temperatures of 6/4 °C and an 8-hour photoperiod to induce dormancy, though the plants did not appear visibly dormant until June 20.

On August 15, the 351 potted cuttings were moved out of dormancy and into a greenhouse where the initial day temperature was 18.5 ± 1.5 °C and night temperature was 16.75 ± 1.25 °C. After the first week, the temperatures were slowly raised to 25.5 ± 2.5 °C during the day and lowered to 10 °C at night. The cuttings were pruned the day they were removed from dormancy so that each cutting had two spurs and each spur had two nodes. Then, the diameter of each spur and node and the distance between the two nodes on each spur were measured with calipers.

Twice a week, beginning August 22, each plant’s development was recorded using the modified Eichorn-Lorenz scale (REFERENCE) and soil moisture was measured with a probe in three locations in each pot. Each spur was kept at two shoots, but only the dominant shoot on each spur had observations recorded. Each shoot was trained up a stake for support. When an inflorescence had developed (EL stage 12), the plant was randomly assigned to one of five growth chambers if it was a part of the heat tolerance experiment. Otherwise, observations on it would continue in the greenhouse.

(NOT SURE WHERE TO PUT THIS) The varieties chosen for inclusion in the experiment expressed a diversity of phenology and had enough reps for at least one plant per chamber.

The five chambers all had a 12-hour photoperiod with 800 m-2s-1 of fluorescent light, but Chamber 1 was set at 17/23 °C, Chamber 2 was set at 23/29 °C, Chamber 3 was set at 27/33 °C, Chamber 4 was set at 31/37 °C, and Chamber 5 was set at 34/40 °C. Initially, CO2 levels were set at 400 ppm during the day and 600 ppm at night, because plants respire at night, increasing CO2 levels (REFERENCE). Each inflorescence was contained in a paper bag to collect the flower caps as they fell.

Observations on the percent flowering, leaf number, stem length, and number fallen flower caps along with soil moisture readings were made three times a week. On September 19, it was noted that some inflorescence bags also contained aborted buds that had yet to flower, and so those numbers were also recorded. Once a plant had reached 100% flowering, or, in the case of plants where the entire inflorescence had died and fallen off, the plant spent a minimum 14 days in the chamber, it was returned to the greenhouse.

All analysis, including analysis of variation (ANOVA) to test for trends between the treatments, was performed in R version 3.3.3 (R Core Team etc). (DOES THIS NEED PROPER CITATION?)